Applicant :Deok-kun Oh et al. Attorney's Docket No.: 21094-002US1 / PO06-0153

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## Amendments to the Claims:

This listing of claims replaces all prior versions and listings of claims in the application:

## Listing of Claims:

1. (Original) A chemically defined medium for fermentation culture of a strain of the genus *Candida*, which comprises 1-10 g/l of urea,1-10g/l of potassium diphosphate, 0.01-1 g/l of magnesium sulfate, 0.1-10 mg/l of MnSO<sub>4</sub> · 4H<sub>2</sub>O, 0.1-10mg/l of CoCl<sub>2</sub> · 6H<sub>2</sub>O, 0.1-10 mg/l of NaMoO<sub>4</sub> · 2H<sub>2</sub>O, 0.1-10 mg/l of ZnSO<sub>4</sub> · 7H<sub>2</sub>O, 0.1-10 mg/l of AlCl<sub>3</sub> · 6H<sub>2</sub>O, 0.1-10 mg/l of CuCl<sub>2</sub> · 2H<sub>2</sub>O, 0.01-5 mg/l of H<sub>3</sub>BO<sub>3</sub>, 1-100 mg/l of FeSO<sub>4</sub> · 7H<sub>2</sub>O, 0.1-10 mg/l of ascorbic acid, 1-100 mg/l of biotin, 1-100 mg/l of choline, and 0.1-10 mg/l of pyridoxine.

2. (Original) A process for producing xylitol in high yield by recycling culture of a strain of the genus *Candida*, which comprises the steps of:

inoculating the strain in a xylose-containing medium and culturing the strain in the xylose-containing medium in a bioreactor;

releasing a culture from the bioreactor and introducing a fresh xylosecontaining medium to the bioreactor continuously;

separating the strain and a culture filtrate from the culture; and recycling the strain to the bioreactor and recovering xylitol from the culture filtrate.

3. (Original The process of claim 2, wherein the strain of the genus *Candida* is *Candida* tropicalis or its mutant strain.

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4. (Currently Amended) The process of claim 2, wherein the xylose-containing medium is [[the]] a chemically defined medium of claim 1 that comprises 1-10 g/l of urea,1-10g/l of potassium diphosphate, 0.01-1 g/l of magnesium sulfate, 0.1-10 mg/l of MnSO<sub>4</sub> · 4H<sub>2</sub>O, 0.1-10 mg/l of CoCl<sub>2</sub> · 6H<sub>2</sub>O, 0.1-10 mg/l of NaMoO<sub>4</sub> · 2H<sub>2</sub>O, 0.1-10 mg/l of ZnSO<sub>4</sub> · 7H<sub>2</sub>O, 0.1-10 mg/l of AlCl<sub>3</sub> · 6H<sub>2</sub>O, 0.1-10 mg/l of CuCl<sub>2</sub> · 2H<sub>2</sub>O, 0.01-5 mg/l of H<sub>3</sub>BO<sub>3</sub>, 1-100 mg/l of FeSO<sub>4</sub> · 7H<sub>2</sub>O, 0.1-10 mg/l of ascorbic acid, 1-100 mg/l of biotin, 1-100 mg/l of choline, and 0.1-10 mg/l of pyridoxine; or a complex medium.

- 5. (Original) The process of claim 2, wherein the culturing is performed by a fed-batch culture or a batch culture.
- 6. (Original) The process of claim 5, wherein in the fed-batch culture, the medium is gradually supplemented with xylose so that the concentration of xylose is maintained at 40-50 g/l on the basis of the medium.
- 7. (Currently Amended) The process of any one of claims 2, 4, 5, and 6 claim 2, wherein the culturing is performed at an agitation speed of 400-600 rpm.
- 8. (Original) The process of claim 2, wherein the separation of the strain and the culture filtrate from the culture is performed by a vacuum microfiltration system or a centrifuge.
- 9. (Currently Amended) The process of claim 2[[ or 8]], wherein the separated strain is concentrated to a density of 10-100 g/l and recycled.
- 10. (New) The process of claim 4, wherein the culturing is performed at an agitation speed of 400-600 rpm.

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11. (New) The process of claim 5, wherein the culturing is performed at an agitation speed of 400-600 rpm.

- 12. (New) The process of claim 6, wherein the culturing is performed at an agitation speed of 400-600 rpm.
- 13. (New) The process of claim 8, wherein the separated strain is concentrated to a density of 10-100 g/l and recycled.